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(54) **OPTICALLY TUNED HARDMASK FOR MULTI-PATTERNING APPLICATIONS**

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See application file for complete search history.

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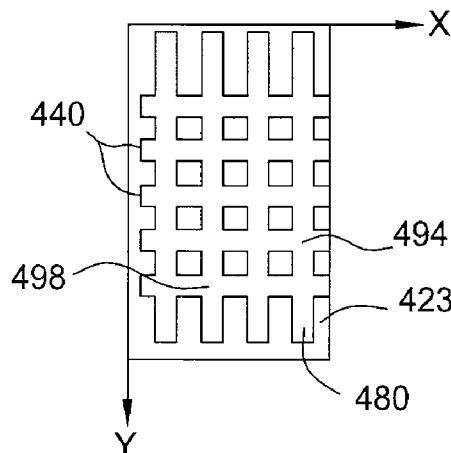
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ABSTRACT

The embodiments herein provides methods for forming a PVD silicon oxide or silicon rich oxide, or PVD SiN or silicon rich SiN, or SiC or silicon rich SiC, or combination of the preceding including a variation which includes controlled doping of hydrogen into the compounds heretofore referred to as $\text{SiO}_x\text{N}_y\text{C}_z\text{H}_w$, where w, x, y, and z can vary in concentration from 0% to 100%, is produced as a hardmask with optical properties that are substantially matched to the photo-resists at the exposure wavelength. Thus making the hardmask optically planarized with respect to the photo-resist. This allows for multiple sequences of litho and etches in the hardmask while the photo-resist maintains essentially no optical topography or reflectivity variations.

13 Claims, 7 Drawing Sheets



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C23C 14/06 (2006.01)
C23C 14/14 (2006.01)
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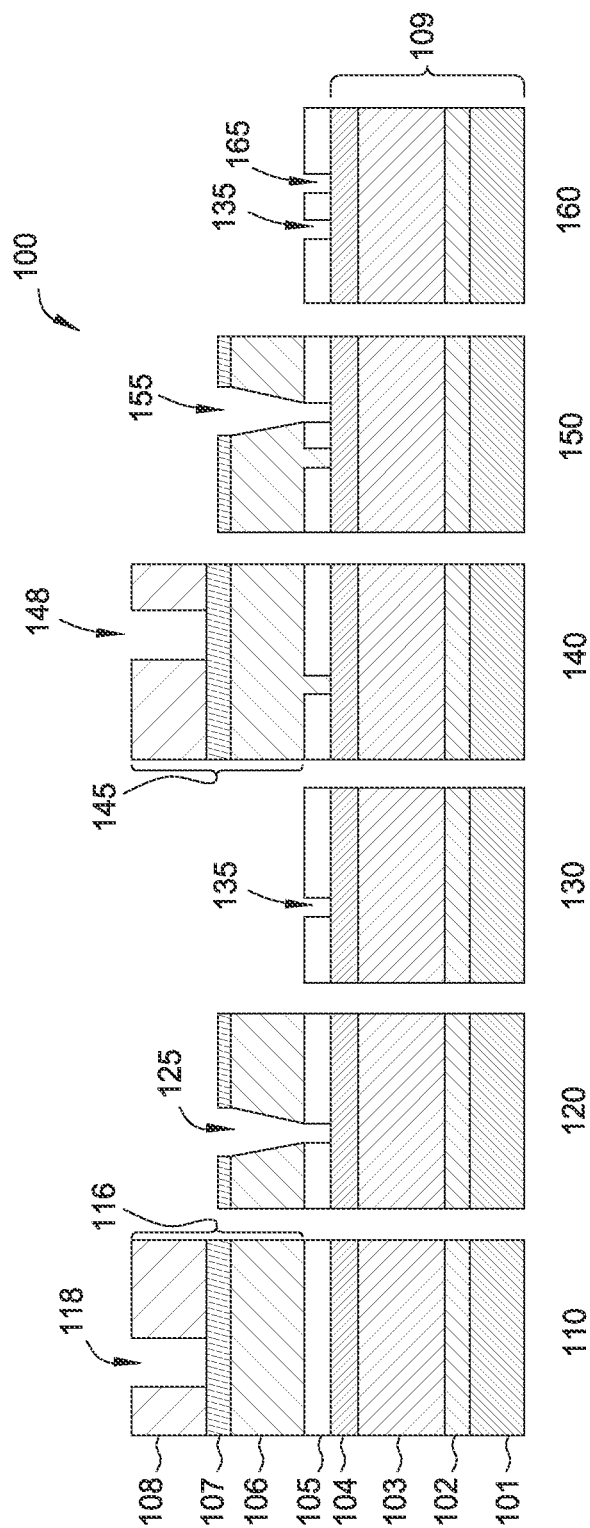


FIG. 1

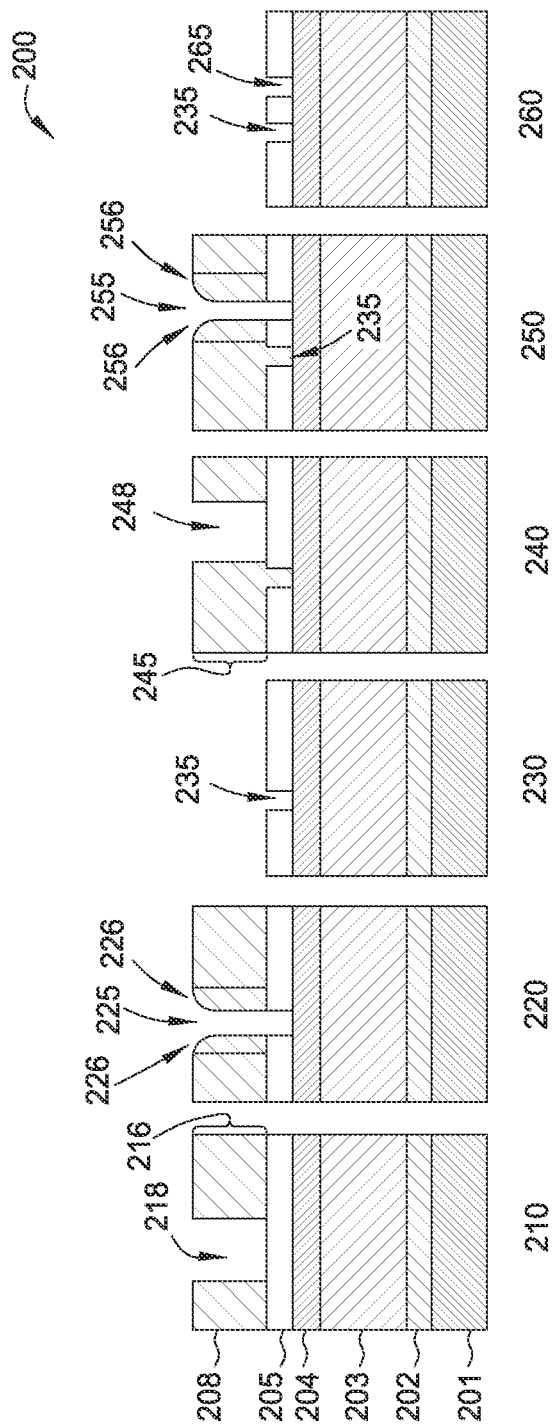
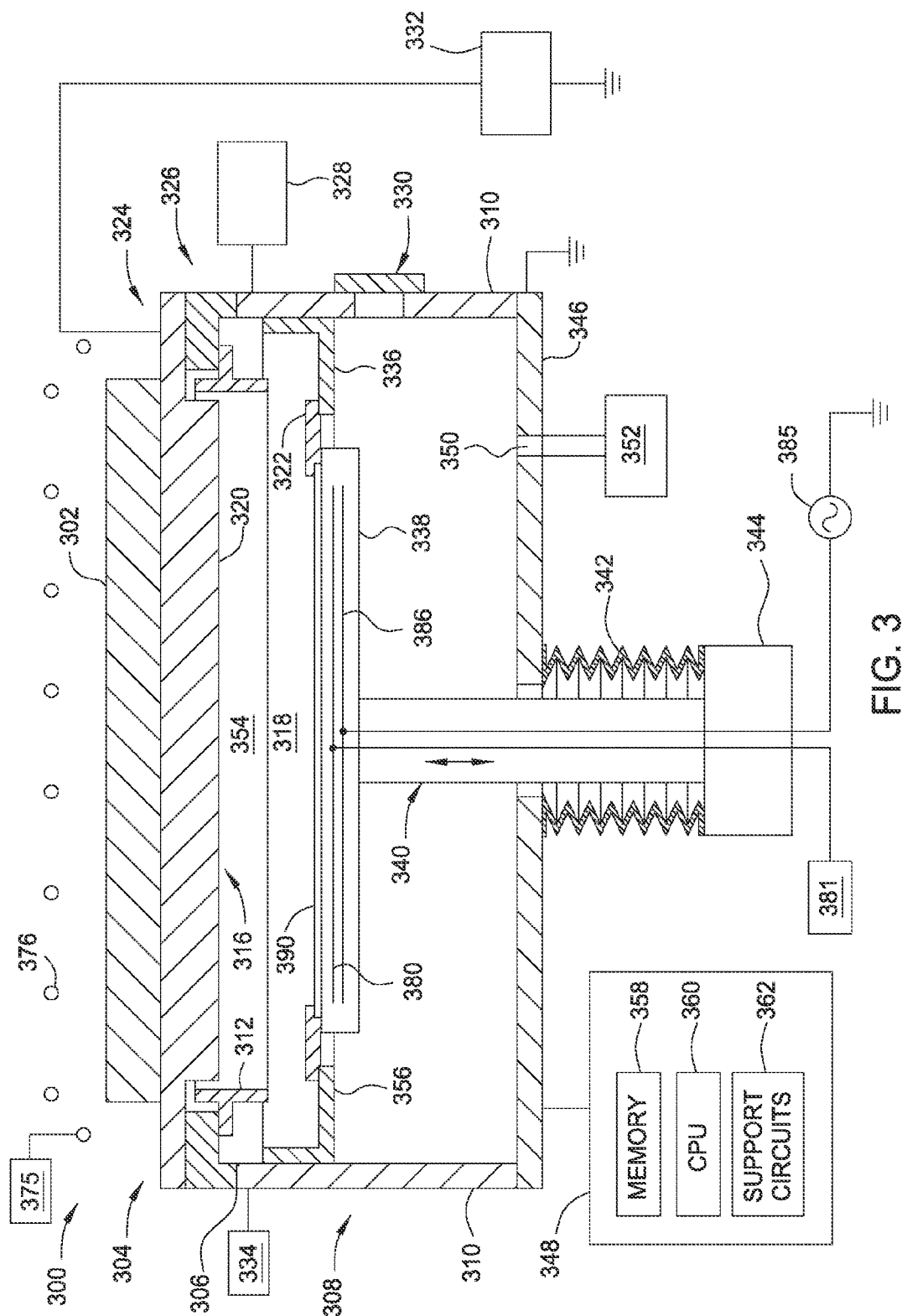
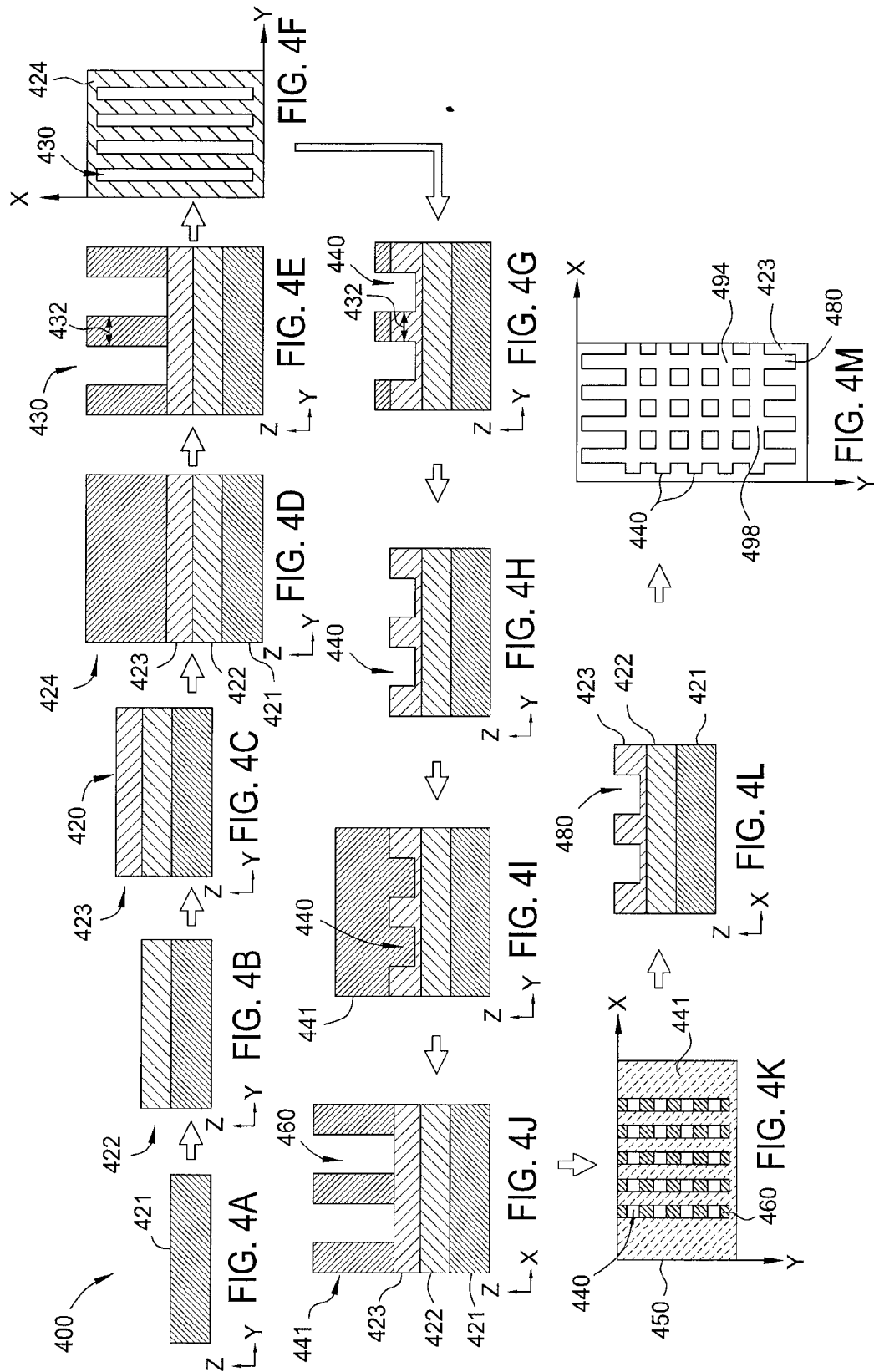


FIG. 2





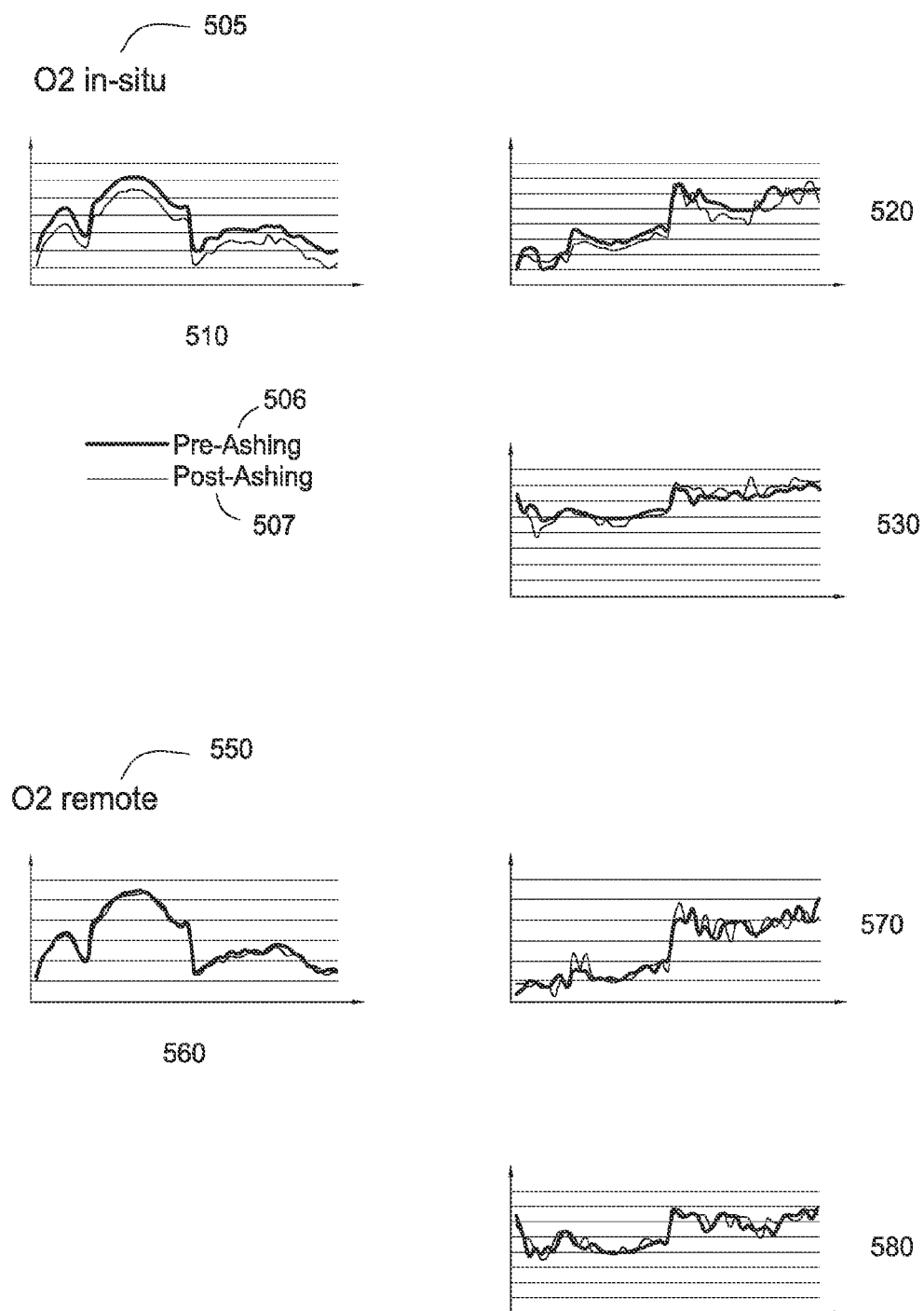


FIG. 5

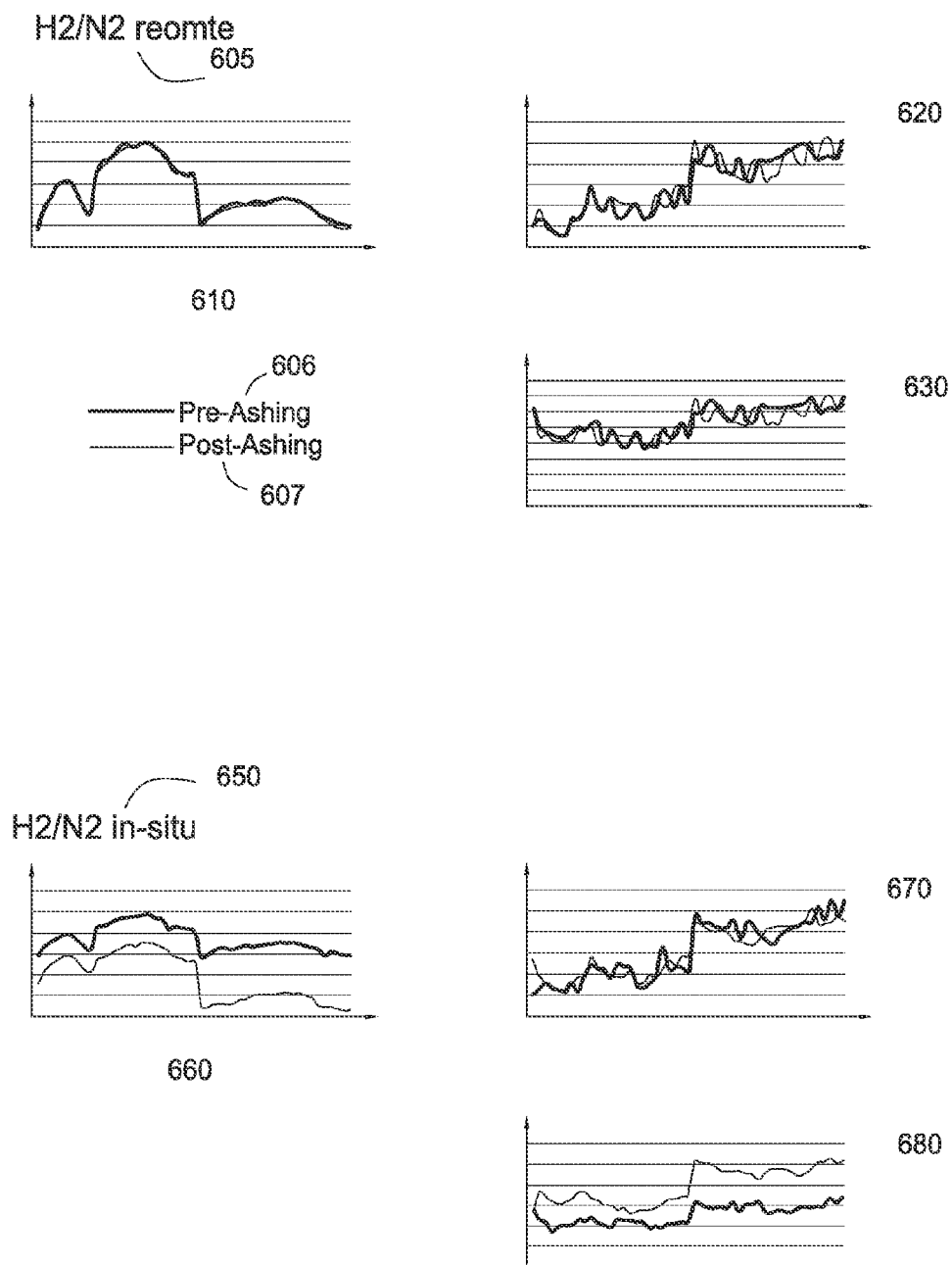


FIG. 6

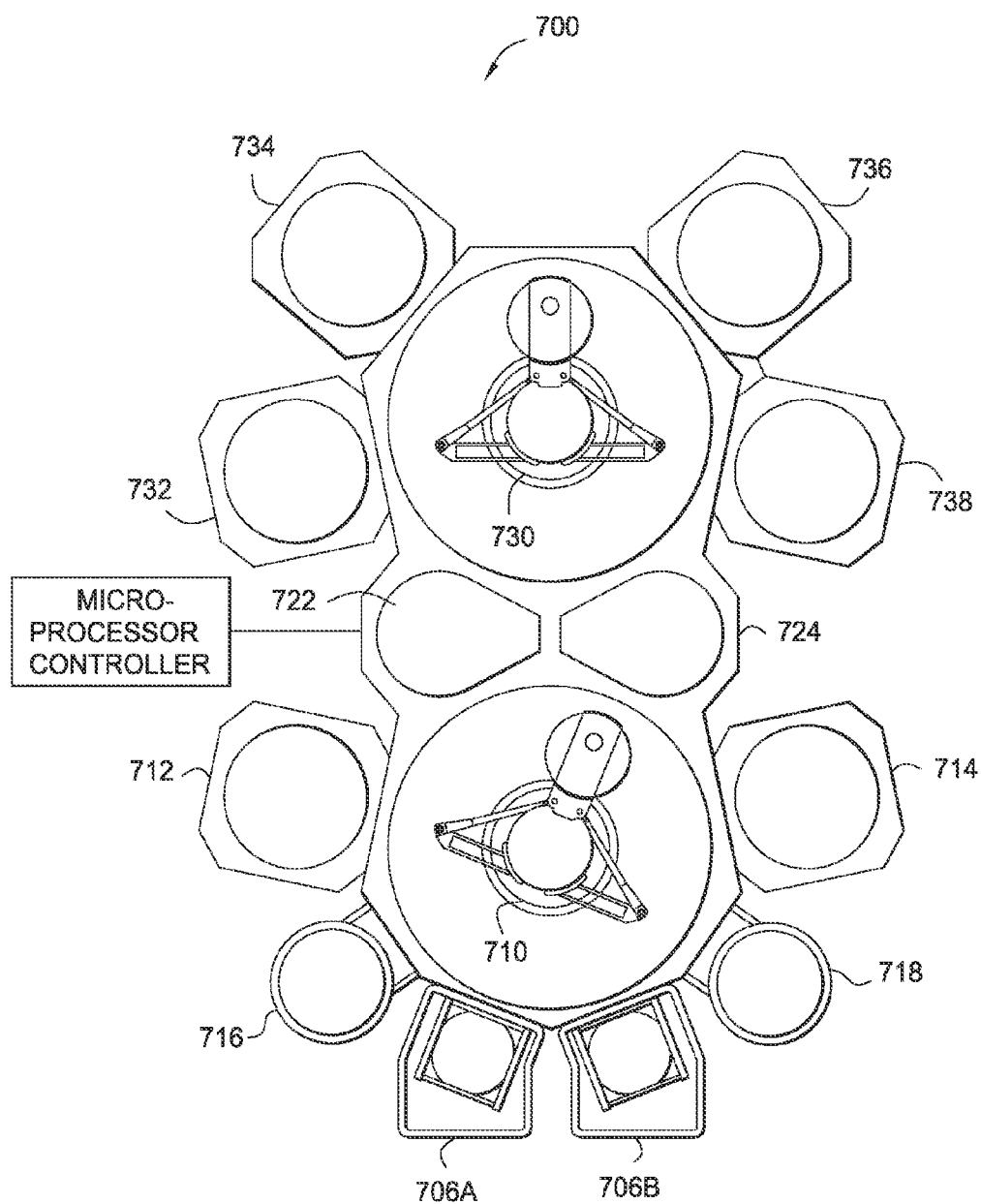


FIG. 7

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OPTICALLY TUNED HARDMASK FOR MULTI-PATTERNING APPLICATIONS

CROSS-REFERENCE TO RELATED APPLICATIONS

This application is a continuation application of U.S. patent application Ser. No. 14/269,010, filed May 2, 2014 and claims benefit of U.S. Provisional Application Ser. No. 61/819,334, filed May 3, 2013, both of which are incorporated by reference in their entirety.

BACKGROUND OF THE DISCLOSURE

1. Field of the Invention

The embodiments herein generally relate to a fabrication process for forming a hardmask for use in a lithographic multi-patterning fabrication process.

2. Description of the Background Art

Reliably producing submicron and smaller features is one of the key requirements of very large scale integration (VLSI) and ultra large scale integration (ULSI) of semiconductor devices. However, with the continued miniaturization of circuit technology, the dimensions of the size and pitch of circuit features, such as interconnects, have placed additional demands on processing capabilities. The multilevel interconnects that lie at the heart of this technology require precise imaging and placement of high aspect ratio features, such as vias and other interconnects. Reliable formation of these interconnects is critical to further increases in device and interconnect density. Additionally, forming sub-micron size features and interconnects with reduced waste of intermediate materials, such as resists and hardmask materials, is desired.

As circuit densities increase for next generation devices, the width or pitch of interconnects, such as vias, trenches, contacts, devices, gates and other features, as well as the dielectric materials there between, are decreasing to 45 nm and 32 nm dimensions. As device scaling was extended to further below the resolution limit of the lithography scanners, multi-patterning was employed to enable meeting the feature density requirements of today's integrated devices. Multi-patterning is a process of performing several resist coating, lithographic patterning, and etching operations to ultimately pattern a film layer in multiple steps. When combined, the overlapping pattern operations form the features in an underlying hardmask layer, which when fully patterned, may be used to pattern an underlying layer, or serve as an implant or diffusion mask.

During simple, non-multi-patterning of an underlying hardmask layer, the current "ultra violet light" wavelengths used for exposure will reflect off the un-patterned interface of the resist and conventional hardmask layer, and may also reflect off the underlying previously formed features, and will as a result, affect the precision of the sidewalls and size of the exposed and developed features in the resist. To correct for this, optical proximity correction (OPC) may be employed in the lithograph mask, which results in an intentional distortion of the position where the resist exposing wavelength reaches the resist, with the result that actually formed developed feature meets a desired feature size and profile. However, with smaller geometries and reflection of the provided exposure's ultra violet electromagnetic energy, OPC is unable to cure distortion effects without additional processing.

One enabler of multiple patterning has been the use of optically opaque films to block the exposure wavelength

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from penetrating to the previously masked hardmask layer, which is sometimes called a memory or memorization, layer. The function of the memory layer is to serve as the hardmask for etching a pattern into a layer there below which may, for example, be a dielectric material, or may, for example, be a mask for another purpose. To pattern the memory layer multiple times, a tri-layer scheme, having an uppermost resist layer, is used in each pattern step. The tri-layer has sufficient opacity to prevent the lithographic resist exposure wavelength from reaching the surface of the memory layer and thus preventing reflections of the exposure electromagnetic energy off of previously formed hardmask features back into the resist, which would result in unintended exposure of areas therein. After each patterning step of the multi-patterning scheme, the tri-layer must be stripped off with wet and/or gas based chemistries, and the wafer and the memory layer must be wet cleaned and dried and a new tri-layer applied thereto before the next pattern of the multi-pattern can be formed in the memory layer.

While the benefits of multiple patterning in terms of resolution, depth of focus and lithographic defect sensitivity are understood, there is additional desire to control the process budget and increase and maintain yield.

Therefore, there is a need for an improved method for lithographically creating a multi-patterned hardmask on a substrate.

SUMMARY OF THE INVENTION

The embodiments herein provide apparatus and methods for performing multi-patterning of a memory or hardmask layer without the need for repeatedly ashing and depositing an optically opaque material or a tri-layer stack, and where only the resist need be stripped, and re-applied after the substrate is cleaned, to perform the next patterning step on the hardmask. In one aspect this is accomplished by applying, as the hardmask layer, a thin film which is optically tuned to match, or very nearly match, the optical characteristics of the resist at the wavelength of the lithography exposure step, thereby providing a hardmask layer which does not create reflection at the resist-hardmask layer interface. In one embodiment, the memory layer is a PVD deposited silicon oxide or silicon rich oxide, or a PVD SiN or silicon rich SiN, or SiC or silicon rich SiC, or a combination of the preceding including a variation which includes controlled doping of hydrogen into the compounds, heretofore referred to as $\text{SiO}_x\text{N}_y\text{C}_z\text{H}_w$, where w, x, y, and z can vary in concentration relative to each other from 0% to 100%. The memory layer, such as the $\text{SiO}_x\text{N}_y\text{C}_z\text{H}_w$ layer, has optical properties that are substantially similar, or nearly matched, to the optical properties of the photo-resist to be formed and patterned thereon, at the exposure wavelength (typically 193 nm for advanced lithography). Thus the interface of the photo-resist and the memory (hardmask) layer is optically "invisible" to the exposure wavelength. This allows for multiple sequences of litho and etching of the hardmask without the need to apply, pattern, and strip intermediate material layers while the exposed photo-resist has essentially no optical topography or reflectivity induced variations from the desired exposure pattern. As a result, each subsequent lithography exposure experiences the same or nearly the same reflectivity, eliminating the need to perform complex optical proximity corrections, and apply multiple complex tri-layers followed by litho, etch and strip of the tri-layer.

The embodiments herein include hardware for forming the optically matched hardmask which includes a pumping

system and chamber cooling system, a full face erosion magnetron cathode, a process kit and gas flow design, an electro-static chuck (ESC), a pulsed DC power supply, a doped silicon target, and H and/or O and/or N and/or C containing gas supplies.

In some embodiments, the hardware is configured to enable formation of an optically matched, to a specifically desired resist, $\text{SiO}_x\text{N}_y\text{C}_z\text{:H}_w$ layer. The $\text{SiO}_x\text{N}_y\text{C}_z\text{:H}_w$ film refractive index (n) and extinction coefficient (K) are tunable by adjusting gas flows and the resultant film w, x, y and z values, to match the optical properties of the film to the resist used to etch the layer.

Embodiments of the disclosure may provide a method for forming a hardmask on a film stack, comprising sputtering a material comprising silicon from a target disposed in a chamber onto a surface of a substrate, and delivering a flow of a process gas while sputtering the material comprising silicon from the target, wherein the process gas comprises oxygen and nitrogen, and wherein the ratio of oxygen to nitrogen in the process gas is adjusted so that an optical property of the sputtered material has an substantially equivalent value as the optical property of a photo-resist layer, which is to be disposed on a surface of the sputtered material, at an intended lithography exposure wavelength.

Embodiments of the disclosure may further provide a hardmask layer, comprising a SiO_xN_y layer disposed on a surface of a substrate, wherein the silicon, oxygen and nitrogen content of SiO_xN_y layer are adjusted so that a refractive index (n) of the SiO_xN_y layer is substantially equal to a refractive index (n) of a resist layer to be formed thereon, and the refractive indices are measured at an intended lithography exposure wavelength. The SiO_xN_y layer may also have an extinction coefficient that is substantially equal to an extinction coefficient of the resist layer to be formed thereon.

BRIEF DESCRIPTION OF THE DRAWINGS

So that the manner in which the above recited features of the embodiments herein are attained and can be understood in detail, a more particular description of the invention, briefly summarized above, may be had by reference to the embodiments thereof which are illustrated in the appended drawings.

FIG. 1 depicts a conventional cycle for multi-patterning a hardmask layer using a tri-Layer etch.

FIG. 2 depicts an embodiment herein for multi-patterning a hardmask layer using a single-layer etch.

FIG. 3 depicts a cross sectional view of one embodiment of a process chamber capable of forming a hardmask layer.

FIGS. 4A to 4M depict a process flow for multi-patterning the hardmask layer using a single-layer etch.

FIG. 5 depicts the changes to a hardmask layer from ashing with in-situ and remote plasma activated O_2 .

FIG. 6 depicts the changes to a hardmask layer from ashing with in-situ and remote plasma activated H_2/N_2 .

FIG. 7 illustrates an exemplary cluster tool 700 suitable for lithographically creating a multi-patterned hardmask on a substrate.

To facilitate understanding of the embodiments, identical reference numerals have been used, where possible, to designate identical elements that are common to the figures. It is contemplated that elements and features of one embodiment may be beneficially incorporated in other embodiments without further recitation.

It is to be noted, however, that the appended drawings illustrate only exemplary embodiments and are therefore not

to be considered limiting of its scope, for the invention may admit to other equally effective embodiments.

DETAILED DESCRIPTION

In some embodiments, a hardmask layer is provided having optical properties, with respect to those of an overlying photo-resist layer, that are substantially similar. The optical properties of the hardmask layer are such that the internal reflections and refractions of light at the photo-resist's lithographic wavelengths do not occur, or are minimized to such an extent that degradation of the precision of the exposed photo-resist feature does not occur. The hardmask layer is provided to be patterned a multiple number of times by repeating the steps of forming a photo-resist on the hardmask layer, exposing the photo-resist, transferring the developed pattern from the photo-resist to the hardmask layer, and stripping/ashing the photo-resist to remove the photo-resist from the hardmask layer and then cleaning and drying the hardmask layer for receipt of another layer of photo-resist directly thereon.

The hardmask layer, also called a memory layer, may be located directly on a semiconductor layer, such as a wafer, and provide a mask thereon for ion implanting or diffusing of dopants into the substrate, and may also be used in the formation of openings in deposited film layers for the formation of interconnects, as well as other features and devices, such as film layers for raised gates, capacitors, etc, which may be formed above the substrate surface.

The embodiment specifically described herein disclose methods for ultimately forming a multi-patterned hardmask ultimately to be used to pattern and etch a dielectric interconnect material for the formation of metal containing features with high aspect ratios and/or with small dimensions therein. As discussed herein, high aspect ratio of the features refers to the submicron structures having aspect ratio in excess of 4:1, and small dimensions refer to the submicron structures having dimensions are about less than 55 nm. The deposition process may include supplying at least an inert gas into the processing chamber during processing. By adjusting the gas ratio and partial pressure of the inert gas supplied in the gas mixture during the deposition process, a good profile control and film uniformity formed across the substrate surface may be obtained.

The embodiments herein provide methods for forming a physical vapor deposition, hereinafter PVD, silicon oxide or silicon rich oxide, or PVD SiN or silicon rich SiN, or SiC or silicon rich SiC, or a combination of the preceding including a variation which includes controlled doping of hydrogen into the compounds, heretofore referred to as $\text{SiO}_x\text{N}_y\text{C}_z\text{:H}_w$, where w, x, y, and z can vary in concentration from 0% to 100%. The $\text{SiO}_x\text{N}_y\text{C}_z\text{:H}_w$ layer is produced as a hardmask with optical properties that are sufficiently matched to a photo-resists that will be applied there over and used to etch the $\text{SiO}_x\text{N}_y\text{C}_z\text{:H}_w$ film layer, which is sometimes broadly referred to herein as a SiONC film layer, at the exposure wavelength (193 nm for advanced lithography), to make the hardmask optically indistinguishable from the photo-resist. In some configurations, the $\text{SiO}_x\text{N}_y\text{C}_z\text{:H}_w$ layer may comprise a $\text{SiO}_x\text{C}_z\text{:H}_w$ layer, a $\text{SiO}_x\text{N}_y\text{:H}_w$ layer or a $\text{SiO}_x\text{N}_y\text{C}_z\text{:H}_w$ layer that has desirable optical properties. The matching of the optical properties of the hardmask and the resist allows for multiple sequences of litho, etch, resist strip and reapplication of resist directly on the hardmask for further hardmask patterning; while the resist remains essentially "optically planarized" with no meaningful optical topography or variations in reflectivity of the resist, for example.

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This enables subsequent lithography exposures to experience the same or substantially the same, reflectivity, eliminating the need to simulate and perform complex optical proximity corrections. Additionally, multiple complex tri-layer cycles of litho, etc and strip, are eliminated, as only a single resist layer need be applied direction on the hardmask.

FIG. 1 depicts an example of a conventional prior art cycle **100** for multi-patterning a hardmask layer using a tri-Layer photo-resist that will be applied thereover and used to etch a SiO₂ film layer. The conventional cycle **100** is depicted in snapshots **110** through **160** of FIG. 1, which are depictions of the same portions of the substrate as processing thereof progresses. In the example, a double hardmask etching layer paradigm is shown, wherein the ultimate layer to be patterned, low-K layer **101**, is covered by a first hardmask **102**, a first optical planarization layer **103** and an ashing barrier **104**, and the memorization layer **105** is formed thereover. Specifically, a substrate **109** includes a low K material **101** over which a titanium nitride (TiN) metal hardmask **102** layer is formed, and the first optical planarization layer **103** and the ashing barrier **104** is formed there over. A memorization layer **105** (a hardmask layer) is disposed on a top surface of the ashing barrier **104**. Disposed on the top of the memorization layer **105** is a tri-layer **116**. The tri-layer includes a second optical planarization layer **106** and a silicon containing anti-reflective layer **107** and a photo-resist **108** formed thereon, in that order. In the prior art, the memorization layer is formed using chemical vapor deposition, wherein silicon and oxygen precursors combine, typically in a plasma environment, to form the hardmask film.

In the first snapshot **110**, lithography was used to expose the photo-resist **108** and a pattern feature **118** was developed in the photo-resist **108**. The feature **118** represents an area where photo-resist **108** was removed by the lithographic exposure and subsequent development of the resist.

In the second snapshot **120**, an etching process is performed to etch a first feature **125** through the second optical planarization layer **106** and the memorization layer **105**. During the etching process, the photo-resist **108** is at least partially etched away and an exposed portion of the silicon anti-reflective layer **107**, the second optical planarization layer **106** and the memorization layer **105** are etched.

In the third snapshot **130**, the tri-layer **116** is ashed or otherwise stripped from the memorization layer **105**. This leaves the partially patterned memorization layer **105** on the top surface of the layer stack. Additionally, the memorization layer has an opening **135** therethrough as a result of the previous etching process as shown in the second snapshot **120**. The opening **135** is only a portion of the total pattern required to be created in the memorization layer **105**. However, due to the created topography, further lithography of the memorization layer **105** requires the re-planarization of, and optical isolation of, the memorization layer **105**. Therefore another tri-layer **145** is formed on top of the memorization layer **105**. This involves depositing a third application of the second optical planarization layer **106** and the silicon anti-reflective layer **107**, followed by the application of the photo-resist **108** there over.

In the fourth snapshot **140**, the newly reformed tri-layer **145** has been patterned after a lithographically exposing and developing the resist layer **108** to form a second pattern **148**. The second pattern **148** is used in the second etch shown in the fifth snapshot **150**. The second etch step removes material (shown as a funnel shaped feature or opening **155**) in the reformed tri-layer **145** and the memorization layer **105**. As shown in a sixth snapshot **160**, after a second stripping

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process, remnants of the tri-layer (**145**) above the memorization layer **105** are removed. The memorization layer **105** now has two formed openings, **135** and **165**.

For each patterning step of the same hardmask layer, the prior tri-layer must be stripped from the substrate, the substrate must be cleaned, and then a new tri-layer must be applied. The operations described in FIG. 1 require the deposition of 6 layers (two tri-layers) to planarize, or optically flatten the wafer in order to maintain focus and to optically isolate the memorization layer **105** for the lithography operation when forming the openings **135** and **165** seen in the sixth snapshot. The embodiments hereinafter discussed provide apparatus and methods for preparing multiple openings in multiple patterning steps in a hardmask layer without the need to optically planarize or isolate the patterned hardmask layer during subsequent lithography steps to further pattern the hardmask layer. FIG. 2 depicts a new cycle **200** for multi-patterning an optically matched hardmask layer, in accordance with one embodiment herein.

The new cycle **200**, as shown in FIG. 2, includes snapshots **210** through **260** for a portion of a substrate by which two openings **235** and **265** are separately formed in the hardmask layer **205** in two different lithography steps. In the example, a substrate includes a low K material **201** having a TiN metal hardmask **202** thereon, and a first optical planarization layer **203** (bottom layer) and an antireflective coating/ashing barrier **204** (middle layer). A “memorization” hardmask layer **205** (a hardmask layer formed thereover) is disposed on a top surface of the ARC/ashing barrier **204**. The ARC/ashing barrier **204** may comprise a thin film for with both antireflective properties (at $\lambda=193$ nm) and etch stopping and ashing barrier properties for the etch and PR strip performed on the hardmask layer **205**. The ARC/ashing barrier **204** may be formed from a Si based material. The ARC/ashing barrier **204** may alternatively be formed from a aluminum nitride (AlN), AlON, SiN, TiN or other robust material having antireflective properties, etch stopping properties, ashing barrier properties and can also be selectively removed. For example, the ARC/ashing barrier **204** may be formed by depositing an AlN layer over the surface of the substrate. The AlN layer may be formed by physical vapor deposition, chemical vapor deposition, atomic layer deposition, or by other suitable method. Disposed on the top of the hardmask layer **205** is only a photo-resist layer **208**, such that the photo-resist layer **208** at least partially is in direct contact with hardmask layer **205**.

The hardmask layer **205** may be a SiO_xN_yC_z:H_w film having a variety of values of w, x, y and z ranging from 0% to 100%. In some cases, the SiON, SiOC or SiONC type films may be doped with hydrogen (H), as noted by the “:H” nomenclature used herein. The hardmask layer **205** composition is tuned to match the n and K values of the photo-resist layer **208** at the exposure wavelength for the lithographic patterning thereof, typically 193 nm. SiO_xN_y films, formed by Physical Vapor Deposition from a silicon target, have a refractive index n ranging between 1.5 to 2.5 and an extinction coefficient K of around 0 to 0.3 at exposure wavelength 193 nm. The termination of the SiO_xN_yC_z:H_w film may affect the properties of next layer formed thereon, i.e. the photo-resist layer **208**. Therefore, in some configurations, it is desirable to adjust the composition and/or properties of the material in the hardmask layer **205** found at the interface of the hardmask layer **205** and the photo-resist layer **208**, or termination region. The termination region of the hardmask layer **205**, which is at the interface of the hardmask layer **205** and the photo-resist layer **208**, may be only a few angstroms or monolayers thick. In one

example, it may be desirable to tune the hydrogen (H) and/or nitrogen (N) concentration in the interface region by turning off the flows of nitrogen and/or hydrogen bearing gas(es) during the latter stages of the deposition process, so that the concentration of H and/or N in the termination region, is at least less than the concentration of H and/or N within rest of the formed layer. In one example, the concentration of hydrogen at the surface of the hardmask layer **205** is less than the average hydrogen concentration through a thickness of the deposited hardmask layer **205** and/or the concentration of nitrogen at the surface of the hardmask layer **205** is less than the average nitrogen concentration through the thickness of the deposited hardmask layer **205**. Bombarding the surface of the deposited hardmask layer **205** with gas atoms, such as ionized argon (Ar) gas, may be used to alter the composition (e.g., H and/or N concentration) or crystal structure of the formed hardmask layer **205**. The bombardment process may be performed by forming a plasma over a surface of the substrate and then biasing the substrate, or substrate support that the substrate is resting on, so that ionized gas atoms in the plasma bombard the surface of the substrate. In one configuration, the bombardment process is performed as a last step in the formation of the hardmask layer **205**, so as to alter and/or adjust the surface energy of the film and wetting angle properties of the surface. Therefore, by use of one of the processes discussed above, the properties of the termination region and surface of the hardmask layer **205** can be adjusted so that the surface is hydrophobic or at least more hydrophobic than a uniform composition type hardmask layer and/or an “as-deposited” hardmask layer. Additionally, by use of at least one of these processes, the hardmask layer **205** may be made inert and stable so as not to poison the photo-resist layer **208** formed thereon. As the stoichiometry of the film is not very dependent on deposition temperature, low temperature deposition is possible.

Additionally, the ions bombarding the surface of the deposited hardmask layer **205** may also be biased to the substrate to promote a smooth surface of the substrate prior to disposing the photo-resist layer **208** thereon. The stronger the bias of the ions to the substrate, the denser and smoother the substrate surface becomes. The bombardment process can be used to smooth the surface of the hardmask layer **205**, so that the surface does not have significant roughness or macroscopic features, such as small bumps or divots, which may scatter the lithography wavelengths when patterning the photo-resist. Advantageously, the pattern in the photo-resist layer **208** is tightly controlled prior to etching critical dimensions in the hardmask layer **205**.

In some embodiments, the properties of the hardmask layer **205** are adjusted so that during the alignment of the lithographic pattern phase of the patterning sequence, the previous alignment marks or alignment patterns formed in the hardmask layer **205** are made distinguishable, at the alignment inspection wavelength(s), from the photo-resist layer disposed over the hardmask layer **205**, which is typically 530 or 630 nm. The refractive index n or extinction coefficient K of the hardmask layer **205** may be different from the photo-resist layer **208** at the alignment wavelength (530 nm/630 nm), while still similar at the lithographic exposure wavelength (193 nm). The extinction coefficient can be sensitive to the wavelength. Therefore, the hardmask layer **205** may be doped with hydrogen (H), boron (B), or potassium (K). The dopant atoms may be added to the target or process gas(es) may be used during the deposition process to adjust the optical properties of the hardmask layer **205** at only the desired wavelength. Therefore the hardmask layer

205 may be tuned to substantially match the refractive index n and extinction coefficient K at the lithographic wavelength of 193 nm, while simultaneously having a different refractive index n or extinction coefficient K at an alignment wavelength of 530 nm or 630 nm. Alternately, a dye may be added to the hardmask layer **205** to make the hardmask layer **205** distinguishable from the photo-resist layer **245** at the lithography wavelength of 193 nm, but not at the alignment wavelength of 530 nm or 630 nm.

As described herein, the optical properties of the hardmask and resist are sufficiently similar to enable the hardmask to repeatedly be patterned using only a single layer of resist thereon. To be optically planar, the reflectivity at the interface of the hardmask and resist need be at most 1-2%. Because the reflectivity is a difference function, one must match n & K to enable no reflection at an interface of two materials; simply matching n is not sufficient. For a matched hardmask layer **205**, relative to the photo-resist layer **208**, the n & K sufficiently match such that the Fresnel equations predict <2% reflectivity. However, in the most advanced lithography, the Fresnel equations should predict <0.5% reflectivity. The difference in the n & K values for a hardmask layer and a photo-resist provide an indicator of a sufficiently match hardmask to a resist. A sufficiently matched hardmask layer **205** to photo-resist layer **208** has a K value within ± 0.01 and an n value within ± 0.01 .

To incorporate nitrogen or hydrogen in the sputtered Si, the level of oxygen present in the sputtering chamber must be very low. Even at O_2/N_2 ratio of 1/10, the “10%” value for O_2 is enough to satisfy, i.e., bond at the exclusion of nitrogen to the dangling Si bonds (bond sites) in the PVD deposited silicon. For example, for 100 atoms of silicon dangling bonds (bond sites), introducing 100 atoms of O_2 and 10,000 atoms of N_2 , the 100 atoms of O_2 will bond with the Si dangling bonds to the exclusion of the N_2 . Thus, little nitrogen will be incorporated into the film. Therefore, to tune the refractive index n by the O_2/N_2 ratio of the film, can be tuned by starving the O_2 gas flow during the deposition process.

In the first snapshot of FIG. 2, 193 nm wavelength “ultra violet light” (electromagnetic energy) is used to expose a pattern onto the photo-resist layer **208**, which is developed to form aperture **218**. The lithography tool simultaneously can print 1 billion such features. However, as chips become more dense, chip designs have called for billions more features. To accomplish this, the lithography tool exposes only a part of the total features to be formed in or on the surface in a single operation. Patterns for the remaining features are aligned on the substrate during subsequent operations. The remaining features are then formed on or in the layer in the subsequent operations. Thus the lithographic exposing of 3 billion features may require at least 3 cycles.

The aperture **218** is part of a larger design pattern to be formed in the hardmask layer **205**. The pattern aperture **218** in the photo-resist layer **208** exposes a top surface of the hardmask layer **205** within the aperture **218**. The hardmask layer **205** has a refractive index (n) and an extinction coefficient (K) matched or nearly matched to the photo-resist layer **208**. By matching or nearly matching the optical properties n and K of the photo-resist layer **208** and hardmask layer **205**, the interface of the photo-resist layer **208** and the hardmask layer **205** does not, or at least only minimally, reflects or refracts the ultra violet light of the lithography wavelength, and thus the interface becomes “invisible” to the lithography exposure wavelength. As a result, the photo-resist layer **208** and the hardmask layer **205** together appear optically flattened or planarized to the ultra

violet exposure wavelength of 193 nm, despite the presence of three-dimensional features at the interface thereof, and, without the need to create an optical planarization layer for each patterning step of an underlying hardmask. In one embodiment the photo-resist layer 208 and the hardmask layer 205 has a refractive index $n=1.6$ and an extinction coefficient $K=0.05$. As a result, the exposure electromagnetic energy will not reflect or refract at the physical interface of the hardmask layer 205 and the overlying photo-resist layer 208.

In the second snapshot 220, an optional conformal polymer 226 was deposited on the exposed surfaces of the photo-resist layer 208 and the hardmask layer 205 and an etching process is performed thereon to create opening 225. 193 nm wavelength lithography has practical limitations on the size (width) of the features, such as trenches, it can form in the photo-resist layer 208. The deposition of the conformal polymer enables narrower features than capable by the 193 nm lithography alone. Alternatively, smaller wavelengths for the lithography may be utilized to create a pattern of the desired width. In such an instance, the conformal polymer would not be used.

In the third snapshot 230, after etching the underlying hardmask layer 205 to form first opening 235, the conformal polymer sidewalls 226 and the photo-resist layer 208 were removed by an ashing operation without significantly changing the refractive index (n) or the extinction coefficient (K) value for the hardmask layer 205. In one embodiment, remote O_2 plasma is used to ash the photo-resist layer 208. Alternatively, a mixture of hydrogen containing gasses and nitrogen containing gasses such as H_2 and N_2 are used to ash the photo-resist layer 208, optionally after being activated in a remote plasma source. Opening 235 represents only a portion of the total pattern required to be created in the hardmask layer 205.

The ashing may be performed with a remote plasma source or in-situ. To support multiple ashing operations, changes in the refractive index (n) and the extinction coefficient (K) value for the hardmask layer 205 must be minimal. FIG. 5 depicts the changes in the hardmask layer 205 for ashing with an O_2 plasma formed in-situ 505 and O_2 plasma formed remotely 550.

The three graphical representations of the effect of O_2 plasma formed in-situ 505, depict memorization layer (i.e., hardmask layer or hardmask layer) thickness across the layer in graph 510, the refractive index of the memorization layer across the layer in graph 520, and the extinction coefficient of the memorization layer across the layer in graph 530. The thickness, refractive index and extinction coefficient, are plotted on the y-axis for graphs 510 through 530. Plotted along the x-axis are 49 (forty nine) measured values at sample positions located along concentric circles from the center to the outside edge of the substrate. The measurements are made pre-ashing 506 and post ashing 507. As shown in the graph 510, the thickness of the memorization layer changes from pre-ashing 506 to post ashing 507. As shown in the graph 520, the refractive index of the memorization layer changes from pre-ashing 506 to post ashing 507. And as shown in the graph 530, the extinction coefficient of the memorization layer changes from pre-ashing 506 to post ashing 507. Ashing using an O_2 plasma formed in-situ showed small thickness change, and a minimal drop in the refractive index in the SiON as the O_2 content increased, which may result by oxidation thereof.

The three graphical representations of the effect of O_2 plasma formed remotely 550 as the ashing medium, depict memorization thickness across the layer in graph 560, the

refractive index of the memorization layer across the layer in graph 570, and the extinction coefficient of the memorization layer across the layer in graph 580. Again, the thickness, refractive index and extinction coefficient, are plotted on the y-axis for graphs 560 through 580. Plotted along the x-axis are 49 measured values at sample positions located along concentric circles from the center to the outside edge of the substrate. As shown in the graph 560, the thickness of the memorization layer is substantially unchanged from pre-ashing 506 to post ashing 507. As shown in the graph 570, the refractive index of the memorization layer is substantially unchanged from pre-ashing 506 to post ashing 507. And as shown in the graph 580, the extinction coefficient of the memorization layer is substantially unchanged from pre-ashing 506 to post ashing 507.

FIG. 6 depicts the changes in the hardmask layer 205 for ashing with a plasma mixture of hydrogen (H_2) containing gasses and nitrogen (N_2) containing gasses (H_2/N_2) formed in-situ 605 and formed remotely 650.

The three graphical representations of the effect of H_2/N_2 plasma formed remotely 605 as the ashing medium, depict memorization thickness across the layer in graph 610, the refractive index of the memorization layer across the layer in graph 620, and the extinction coefficient of the memorization layer across the layer in graph 630 is the same manner as FIG. 5. The thickness, refractive index and extinction coefficient, are plotted on the y-axis for graphs 610 through 630. Plotted along the x-axis are 49 measured values at sample positions located along concentric circles from the center to the outside edge of the substrate. The measurements are made pre-ashing 606 and post ashing 607. As shown in the graph 610, the thickness of the memorization layer is substantially unchanged from pre-ashing 606 to post ashing 607. As shown in the graph 620, the refractive index of the memorization layer is substantially unchanged from pre-ashing 606 to post ashing 607. And as shown in the graph 630, the extinction coefficient of the memorization layer is substantially unchanged from pre-ashing 606 to post ashing 607.

The three graphical representations of the effect of H_2/N_2 plasma formed in-situ 650, depict memorization thickness across the layer in graph 660, the refractive index of the memorization layer across the layer in graph 670, and the extinction coefficient of the memorization layer across the layer in graph 680. Again, the thickness, refractive index and extinction coefficient, are plotted on the y-axis for graphs 660 through 680. Plotted along the x-axis are 49 measured values at sample positions located along concentric circles from the center to the outside edge of the substrate. As shown in the graph 660, the thickness of the memorization layer changes from pre-ashing 606 to post ashing 607. As shown in the graph 670, the refractive index of the memorization layer changes from pre-ashing 606 to post ashing 607. And as shown in the graph 680, the extinction coefficient of the memorization layer changes from pre-ashing 606 to post ashing 607. Ashing using an H_2/N_2 plasma formed in-situ showed a certain amount of increase in the extinction coefficient.

Therefore for an O_2 gas, or an H_2/N_2 gas, used as an ashing medium, forming a plasma remotely for ashing the resist from the memorization layer (hardmask layer) 205, causes significantly less change in the refractive index (n) and the extinction coefficient (K) value of the hardmask layer 205 than using a plasma formed in-situ. This allows for multiple sequences of lithography, etch, and strip of a resist to form features in a hardmask, without suffering optical topography or reflectivity variations.

Referring again to FIG. 2 at snapshot 240, to enable a second patterning of the hardmask layer 205, a photo-resist 245 is deposited and developed on the hardmask layer 205 for the purpose of patterning the remaining openings in the hardmask layer 205. Prior to each lithography operation, the patterns of the openings are aligned on the substrate with the patterns etched in the hardmask layer 205 using a 530 nm or 630 nm wavelength. The hardmask layer 205 has a refractive index (n) and an extinction coefficient (K) matching the photo-resist layer 245 at the lithography wavelength of 193 nm, the interface of the hardmask layer 205 and the photo-resist layer 245 is optically indistinguishable to the exposure electromagnetic energy. However, the hardmask layer 205 has a refractive index (n) and an extinction coefficient (K) which is different from the photo-resist layer 245 at a wavelength of about 530 nm or about 630 nm to allow the alignment of the lithography pattern. Thus, the n and K of the hardmask layer 205 is indistinguishable from the n and K of the photo-resist 245 at the wavelength of 193 nm while distinguishable at the wavelength of 530 nm or 630 nm. This removes the need to deposit additional material on the hardmask layer 205 as was previously required to optically planarize or shield the previously etched features from exposure to the lithography electromagnetic energy. That is, the deposition of the tri-layer containing the second optical planarization layer 106 and the silicon anti-reflective layer 107, as shown in FIG. 1, is no longer required. Additionally, since there are no, or acceptably minimal, interface reflections between the hardmask layer 205 and the overlying resist at the lithographic wavelength of 193 nm, optical proximity correction may not be needed for correcting the topology in a following lithographic operation, and applicants have accomplished patterning with the optically matched resist and memorization layer without using optical proximity correction.

In the fourth snapshot 240, a pattern 248 has been lithographically exposed and developed in the photo-resist 245. The pattern 248 is part of a larger design pattern to be formed in the hardmask layer 205. A conformal polymer is deposited on the exposed surfaces of the photo-resist layer 208 and the hardmask layer 205 prior to an etch process. The deposition of the conformal polymer may alternatively occur during the etching of the hardmask layer 205.

In the fifth snapshot 250, an opening 265 in the hardmask layer 205 is shown after an etch operation. After etching, the conformal polymer is visible on the sidewalls 256 of the pattern 248. In a sixth snapshot 260, the conformal polymer sidewalls 256 and the photo-resist layer 245 were removed by an ashing operation. The ashing operation is performed in a manner which does not meaningfully change the refractive index (n) or the extinction coefficient (K) value for the hardmask layer 205. In an integrated circuit fabrication process, substrates may be lithographically processed multiple times. However, the multiple lithographic cycles of etching and ashing performed on the same hardmask layer 205 does not significantly alter the refractive index (n) or the extinction coefficient (K) value of the hardmask layer 205. As a result, the need to deposit an optical planarization layer and a silicon anti-reflective layer and use optical proximity corrections to form openings 235, 265 is negated.

Through matching both the n (refractive index) and the K value for the hardmask to the photo-resists at the 193 nm wavelength, a hardmask, $\text{SiO}_x\text{N}_y\text{C}_z$, is formed. In another embodiment, the hardmask is SiO_2 . Again, this involves tuning the n and K in the hardmask to match the n and K of the photo-resist (soft mask) at the 193 nm wavelength while distinguishing them at the 530 nm or 630 nm wavelength.

The prior art chemical vapor deposition (CVD) process used to form SiON film layers requires higher temperatures, and yields a film having a higher K value because hydrogen becomes incorporated into the film from the Silane precursor. Additionally, the prior art film layers typically contain N—H or amine bonds that form acids which interfere with the photo-resist and may cause byproducts, i.e. scumming, which may need to be removed. The hydrogen forms Si—H which absorbs the 193 nm wavelength and therefore increases the K value. This is especially true at the low deposition temperatures required to meet the thermal budget of the SOC film (typically <200 C). Thus, using CVD, the extinction coefficient (K), or the absorption of electromagnetic energy, of the SiON layer becomes too high. Applicants disclose herein that a Physical Vapor deposition (PVD) formed $\text{SiO}_x\text{N}_y\text{C}_z$ film, or layer, may be formed which is tunable to the n and K values of the relevant resist. Additionally, the hardmask layer may be formed well below 200 C, and acceptable optically tuned hardmask layers have been deposited around 20 to 25 C by sputtering silicon in a reactive atmosphere.

However, silicon is difficult to sputter. Usually, when sputtering a dielectric layer, the surface will accumulate charges leading up to arcing and the ejection of particles of the target. The use of a pulse DC allows fast switching from negative, for sputtering, to positive, for sweeping or charge scrubbing, (neutralizing all the charges on the dielectric surface) of the target.

FIG. 3 illustrates an exemplary physical vapor deposition (PVD) process chamber 300 (e.g., a sputter process chamber) suitable for sputter depositing materials. One example of the process chamber that may be adapted to form the $\text{SiO}_x\text{N}_y\text{C}_z$ film layer is a PVD process chamber, available from Applied Materials, Inc., located in Santa Clara, Calif. It is contemplated that other sputter process chambers, including those from other manufactures, may be adapted to practice the present invention.

The process chamber 300 includes a chamber body 308 having a processing volume 318 defined therein. The chamber body 308 has sidewalls 310 and a bottom 346. The dimensions of the chamber body 308 and related components of the process chamber 300 are not limited and generally are proportionally larger than the size of the substrate 390 to be processed. Any suitable substrate size may be processed. Examples of suitable substrate sizes include substrate with 200 mm diameter, 300 mm diameter or 450 mm diameter.

A chamber lid assembly 304 is mounted on the top of the chamber body 308. The chamber body 308 may be fabricated from aluminum or other suitable materials. A substrate access port 330 is formed through the sidewall 310 of the chamber body 308, facilitating the transfer of a substrate 390 into and out of the process chamber 300. The access port 330 may be coupled to a transfer chamber and/or other chambers of a substrate processing system.

A water vapor generating (WVG) system 334 is coupled to the processing volume 318 defined in the process chamber 300. The WVG system 334 generates ultra-high purity water vapor by means of a catalytic reaction of O_2 and H_2 . Alternatively, the WVG system 334 may also generate water vapor by directly vaporizing water (H_2O) into water vapor as needed. The WVG system 334 has a catalyst-lined reactor or a catalyst cartridge in which water vapor is generated by means of a chemical reaction. The catalyst may include a metal or alloy, such as palladium, platinum, nickel, combinations thereof and alloys thereof. While water vapor is usually generated by flowing H_2 and O_2 into the reactor, the

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O₂ may be supplemented or substituted with another oxygen source compound, such as NO, N₂O, NO₂, N₂O₅, H₂O₂ or O₃. In one embodiment, H₂ and N₂O are utilized to form a water vapor as needed. The ultra-high purity water is ideal for providing H₂O into the processing volume **318** for forming plasma having O₂ and H₂ ions.

A gas source **328** is coupled to the chamber body **308** to supply process gases into the processing volume **318**. In one embodiment, process gases may include inert gases, non-reactive gases, and reactive gases if necessary. Examples of process gases that may be provided by the gas source **328** include, but not limited to, argon gas (Ar), helium (He), neon gas (Ne), krypton (Kr), xenon (Xe), nitrogen gas (N₂), oxygen gas (O₂), hydrogen gas (H₂), H₂O (in vapor form from the WVG system **334**), forming gas (N₂+H₂), ammonia (NH₃), methane (CH₄), carbon monoxide (CO), and/or carbon dioxide (CO₂), among others.

A pumping port **350** is formed through the bottom **346** of the chamber body **308**. A pumping device **352** is coupled to the processing volume **318** to evacuate and control the pressure therein. A pumping system and chamber cooling design enables high base vacuum (1E-8 Torr or less) and low rate-of-rise (1,000 mTorr/min) at temperatures suited to thermal budget needs, e.g., -25 C to +500 C. The pumping system is designed to provide precise control of process pressure which is a critical parameter for RI control and tuning.

The lid assembly **304** generally includes a target **320** and a ground shield assembly **326** coupled thereto. The target **320** provides a material source that can be sputtered and deposited onto the surface of the substrate **390** during a PVD process. Target **320** serves as the cathode of the plasma circuit during DC sputtering.

The target **320** or target plate may be fabricated from a material utilized for the deposition layer, or elements of the deposition layer to be formed in the chamber. A high voltage power supply, such as a power source **332**, is connected to the target **320** to facilitate sputtering materials from the target **320**. In one embodiment, the target **320** may be fabricated from a material containing silicon (Si), titanium (Ti) metal, tantalum metal (Ta), hafnium (Hf), tungsten (W) metal, cobalt (Co), nickel (Ni), copper (Cu), aluminum (Al), alloys thereof, combinations thereof, or the like. Additionally, the electron emission from the target during processing may be controlled by n-type or p-type doping of the target. The target may be doped with a conducting element such as Boron (B). The target material may have a monocrystalline versus a polycrystalline structure. For example, the target may include Si in which the crystal lattice of the entire Si target is a single crystal. In an exemplary embodiment depicted herein, the target may be fabricated from an intrinsic Si, or doped electrically conductive Si, or specific composition SiO_xN_yC_zH_w composite target. In one embodiment, the target is 99.999% pure Si doped with Boron to about 1×10¹⁸ atoms per cm².

The target **320** generally includes a peripheral portion **324** and a central portion **316**. The peripheral portion **324** is disposed over the sidewalls **310** of the chamber. The central portion **316** of the target **320** may have a curvature surface slightly extending towards the surface of the substrate **390** disposed on a substrate support **338**. The spacing between the target **320** and the substrate support **338** is maintained between about 50 mm and about 350 mm. It is noted that the dimension, shape, materials, configuration and diameter of the target **320** may be varied for specific process or substrate requirements. In one embodiment, the target **320** may further include a backing plate having a central portion bonded

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and/or fabricated by a material desired to be sputtered onto the substrate surface. The target **320** may also include adjacent tiles or segmented materials that together form the target.

The lid assembly **304** may further comprise a full face erosion magnetron cathode **302** mounted above the target **320** which enhances efficient sputtering materials from the target **320** during processing. The full face erosion magnetron cathode **321** allows easy and fast process control and tailored film properties while ensuring consistent target erosion and uniform deposition of SiO_xN_yC_zH_w film across the wafer for a variety of values of w, x, y and z ranging from 0% to 100%. Examples of the magnetron assembly include a linear magnetron, a serpentine magnetron, a spiral magnetron, a double-digitated magnetron, a rectangularized spiral magnetron, among others.

The ground shield assembly **326** of the lid assembly **304** includes a ground frame **306** and a ground shield **312**. The ground shield assembly **326** may also include other chamber shield member, target shield member, dark space shield, and dark space shield frame. The ground shield **312** is coupled to the peripheral portion **324** by the ground frame **306** defining an upper processing region **354** below the central portion of the target **320** in the processing volume **318**. The ground frame **306** electrically insulates the ground shield **312** from the target **320** while providing a ground path to the chamber body **308** of the process chamber **300** through the sidewalls **310**. The ground shield **312** constrains plasma generated during processing within the upper processing region **354** and dislodges target source material from the confined central portion **316** of the target **320**, thereby allowing the dislodged target source to be mainly deposited on the substrate surface rather than chamber sidewalls **310**. In one embodiment, the ground shield **312** may be formed by one or more work-piece fragments and/or a number of these pieces bonding by processes known in the art, such as welding, gluing, high pressure compression, etc.

A shaft **340** extending through the bottom **346** of the chamber body **308** couples to a lift mechanism **344**. The lift mechanism **344** is configured to move the substrate support **338** between a lower transfer position and an upper processing position. A bellows **342** circumscribes the shaft **340** and coupled to the substrate support **338** to provide a flexible seal there between, thereby maintaining vacuum integrity of the chamber processing volume **318**.

The substrate support **338** may be an electro-static chuck and have an electrode **380**. The electro-static chuck (ESC) **338** uses the attraction of opposite charges to hold both insulating and conducting substrates **390** for a lithography processes and is powered by DC power supply **381**. The ESC **338** comprises an electrode embedded within a dielectric body. The DC power supply **381** may provide a DC chucking voltage of about 200 to about 2000 volts to the electrode. The DC power supply **381** may also include a system controller for controlling the operation of the electrode **380** by directing a DC current to the electrode for chucking and de-chucking the substrate **390**.

The temperature of the PVD process may be kept below the temperature at which organic films deposited beneath the hardmask layer **205** become volatile. For example, temperature may be less than about 250 Celsius and have about a 50 Celsius margin to prevent the organic films below the hardmask layer **205** from gassing out and contaminating the chamber. The ESC **338**. The ESC **338** performs in the temperature range required by the thermal budget of the device integration requirements. For example, Detachable ESC **338** (DTEESC) for minus 25 C to 100 C temperature

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range, Mid-Temp ESC **338** (MTEC) for 100 C to 200 C temperature range, High Temperature or High Temperature Biasable or High Temperature High Uniformity ESC **338** (HTESC or HTBESC or HTHUESC) for temperatures ranging from 200 C to 500 C that ensures fast and uniform heating up of wafers.

After the process gas is introduced into the PVD chamber **300**, the gas is energized to form plasma. An antenna **376**, such as one or more inductor coils, may be provided adjacent the PVD chamber **300**. An antenna power supply **375** may power the antenna **376** to inductively couple energy, such as RF energy, to the process gas to form plasma in a process zone in the PVD chamber **300**. Alternatively, or in addition, process electrodes comprising a cathode below the substrate **390** and an anode above the substrate **390** may be used to couple RF power to generate plasma. A controller that also controls the operation of other components in the PVD chamber **300** may control the operation of the power supply **375**.

A shadow frame **322** is disposed on the periphery region of the substrate support **338** and is configured to confine deposition of source material sputtered from the target **320** to a desired portion of the substrate surface. A chamber shield **336** may be disposed on the inner wall of the chamber body **308** and have a lip **356** extending inward to the processing volume **318** configured to support the shadow frame **322** disposed around the substrate support **338**. As the substrate support **338** is raised to the upper position for processing, an outer edge of the substrate **314** disposed on the substrate support **338** is engaged by the shadow frame **322** and the shadow frame **322** is lifted up and spaced away from the chamber shield **336**. When the substrate support **338** is lowered to the transfer position adjacent to the substrate transfer access port **330**, the shadow frame **322** is set back on the chamber shield **336**. Lift pins (not shown) are selectively moved through the substrate support **338** to lift the substrate **390** above the substrate support **338** to facilitate access to the substrate **390** by a transfer robot or other suitable transfer mechanism.

A controller **348** is coupled to the process chamber **300**. The controller **348** includes a central processing unit (CPU) **360**, a memory **358**, and support circuits **362**. The controller **348** is utilized to control the process sequence, regulating the gas flows from the gas source **328** into the process chamber **300** and controlling ion bombardment of the target **320**. The CPU **360** may be of any form of a general purpose computer processor that can be used in an industrial setting. The software routines can be stored in the memory **358**, such as random access memory, read only memory, floppy or hard disk drive, or other form of digital storage. The support circuits **362** are conventionally coupled to the CPU **360** and may comprise cache, clock circuits, input/output subsystems, power supplies, and the like. The software routines, when executed by the CPU **360**, transform the CPU into a specific purpose computer (controller) **348** that controls the process chamber **300** such that the processes are performed in accordance with the present invention. The software routines may also be stored and/or executed by a second controller (not shown) that is located remotely from the PVD chamber **300**.

During processing, material is sputtered from the target **320** and deposited on the surface of the substrate **390**. The target **320** and the substrate support **338** are biased relative to each other, and/or relative to ground, by the power source **332** to maintain a plasma formed from the process gases supplied by the gas source **328**. The ions from the plasma are accelerated toward and strike the target **320**, causing target

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material to be dislodged from the target **320**. The dislodged target material and reactive process gases together form a layer on the substrate **390** with desired compositions. RF, DC or fast switching pulsed DC power supplies or combinations thereof provide tunable target bias for precise control of sputtering composition and deposition rates for the $\text{SiO}_x\text{N}_y\text{C}_z\text{:H}_w$ material.

In some embodiments, it is also desirable to separately apply a bias to the substrate during different phases of the $\text{SiO}_x\text{N}_y\text{C}_z\text{:H}_w$ layer deposition process. Therefore, a bias may be provided to a bias electrode **386** (or chuck electrode **380**) in the substrate support **338** from a source **385** (e.g., DC and/or RF source), so that the substrate **390** will be bombarded with ions formed in the plasma during one or more phase of the deposition process. Biasing the electrode can be used to smooth the surface of the substrate and increase the hydrophobicity for the surface of the substrate. In some process examples, the bias is applied to the substrate after the $\text{SiO}_x\text{N}_y\text{C}_z\text{:H}_w$ film deposition process has been performed. Alternately, in some process examples, the bias is applied during the $\text{SiO}_x\text{N}_y\text{C}_z\text{:H}_w$ film deposition process. Thus, when the substrate bias is maintained throughout the deposition process the bombarding atoms add kinetic energy to the depositing material found at the surface of the substrate. For example, between about 50 Watts and about 1100 Watts of energy may be used to bias ions to the substrate to form a smooth dense film. A larger bias drives the ions with greater energy to the substrate surface. For example, a post treatment process may use an argon (Ar) containing gas and smooth the substrate film by providing bias energy, such between about 200-1100 Watts, to direct the ions at the substrate surface of the $\text{SiO}_x\text{N}_y\text{C}_z\text{:H}_w$ film, i.e. the hardmask layer **205**. The smooth film surface for the hardmask layer advantageously prevents light scattering on bumpy surfaces for the lithography which may affect the quality of the features formed in the photo-resist layer. In another example, a post treatment process may use a process gas that includes argon (Ar) gas. The post treatment may make the surface of the $\text{SiO}_x\text{N}_y\text{C}_z\text{:H}_w$ film more stable and hydrophobic.

The combination of the unique hardware and process produces an optically matched, to a resist to be formed thereon, $\text{SiO}_x\text{N}_y\text{C}_z\text{:H}_w$. The $\text{SiO}_x\text{N}_y\text{C}_z\text{:H}_w$ film refractive index (n) and extinction coefficient (K) are tunable by adjusting gas flows and the resultant film stoichiometry. A target is poisoned when the surface is no longer conductive. So, poison mode sputtering occurs when the reactive species in the gas atmosphere at the target surface is specifically rich in reactive species that a dielectric layer is formed across the face of the target and the target behaves electrically like a dielectric and not a conductor or semiconductor. H and/or O and/or N and/or C containing gases can react on and poison the surface of the target to create the desired $\text{SiO}_x\text{N}_y\text{C}_z\text{:H}_w$ stoichiometry to enable the combination of optical "invisibility" and etch selectivity best suited for the device needs. In addition, methane (CH_4), carbon monoxide (CO), hydrogen (H_2), or carbon dioxide (CO_2) can also be used as a poisoning gas to achieve more tunability of the n and K values along with a desired etch selectivity. For example, H_2 may be used to form a thin film or a SiH monolayer in order to increase the light absorption K without increasing the reflectivity n due to the layer tenuity. Thus, the H containing gas may form a light absorption layer.

The hardware may also include a gas box. Mixing gas in a gas box before coming into the chamber is one way to ensure a homogenous gas mixture of the various process gases for uniform target poisoning. The gas box would mix

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H₂ with N₂ for fine tuning the K . O₂ would be added to the NH mixture (more stable and less dangerous than mixing O₂ and H₂ first). Two O₂ mass flow controllers (MFC's) are utilized. One for bulk control of the O₂, for around 100-200 sccm, and one for fine control of the O₂, for around 5 sccm. Process argon (Ar) could also be added to the mixture. This ensures a homogenous gas mixture before it enters the chamber. In one embodiment, The H and N gas may be turned off prior to the complete formation of the SiO_xN_yC_z:H_w film so as to not poison the photo-resist layer to be formed on top. In one example, a gas flow mixture ratio of 20% O₂/60% N₂/20% Ar is provided to a processing region of a chamber that is maintained at 3.5 mTorr while a 3 kW pulsed DC power is delivered to the target. However, in some cases, the gas flow mixture may be provided to the processing region of the chamber while a DC, a pulsed DC, an RF and/or a pulsed RF power is delivered to the target.

The exemplary physical vapor deposition (PVD) process chamber 300 may be part of a cluster tool. FIG. 7 illustrates an exemplary cluster tool 700 suitable for creating and etching a multi-patterned hardmask on a substrate. The cluster tool 700 features at least one physical vapor deposition (PVD) chamber 300, as described above. An example of the cluster tool 700 is the Endura® system available from Applied Materials, Inc., of Santa Clara, Calif. Cluster tools manufactured by others may be used as well.

The cluster tool 700 can include one or more load lock chambers 706A, 706B for transferring of substrates into and out of the cluster tool 700. Typically, since the cluster tool 700 is under vacuum, the load lock chambers 706A, 706B may "pump down" the substrates introduced into the cluster tool 700. A first robot 710 may transfer the substrates between the load lock chambers 706A, 706B, and a first set of one or more substrate processing chambers 712, 714, 716, 718 (four are shown). Each processing chamber 712, 714, 716, 718, can be outfitted to perform a number of substrate processing operations including the etch processes described herein in addition to cyclical layer deposition (CLD), atomic layer deposition (ALD), chemical vapor deposition (CVD), physical vapor deposition (PVD) such as PVD chamber 300, pre-clean, degas, orientation and other substrate processes.

The first robot 710 can also transfer substrates to/from one or more intermediate transfer chambers 722, 724. The intermediate transfer chambers 722, 724 can be used to maintain ultrahigh vacuum conditions while allowing substrates to be transferred within the cluster tool 700. A second robot 730 can transfer the substrates between the intermediate transfer chambers 722, 724 and a second set of one or more processing chambers 732, 734, 736, 738. Similar to processing chambers 712, 714, 716, 718, the processing chambers 732, 734, 736, 738 can be outfitted to perform a variety of substrate processing operations including the etch processes described herein in addition to cyclical layer deposition (CLD), atomic layer deposition (ALD), chemical vapor deposition (CVD), physical vapor deposition (PVD), pre-clean, thermal process/degas, and orientation, for example. Any of the substrate processing chambers 712, 714, 716, 718, 732, 734, 736, 738 may be removed from the cluster tool 700 if not necessary for a particular process to be performed by the cluster tool 700.

An illustrative multi-processing cluster tool 700 for forming the structures of FIGS. 4A-4M can include up to four PVD chambers 732, 734, 736, 738 (with an option for a fifth chamber between 734 and 736) similarly configured to the PVD chamber 300 described above. The PVD or ALD chambers 712 or 714 may be configured to deposit a thin ARC/ashing layer (e.g., AlN or SiN or TiN). The thermal

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treatment chambers 716, 718 may be capable of performing a thermal treatment process, such as a degas process. The degas process may remove potential contaminants which may otherwise outgas during chamber operations. In one configuration the thermal treatment chamber 716, 718 is adapted to complete a degas process at a temperature of about 200-300 degrees C. and pressure in the 100 mTorr or less range. Additionally, the thermal treatment chamber 716, 718 may preheat the substrates prior to processing in the cluster tool 700.

The cluster tool 700 may be used to performed the methods described in FIG. 2 above and FIG. 4 below. During processing, a substrate that is to be processed may arrive to the cluster tool 700 in a pod (not shown). The substrate is transferred from the pod to the vacuum compatible load lock 706A, 706B by the factory interface robot (not shown). The first robot 710 then moves the substrate into the degas chamber 716 or 718 for degassing and preheating. The first robot 710 then picks up the substrate from the degas chamber 716 or 718 and loads the substrate into the intermediate transfer chamber 722 or optionally into chamber 712 or 714 for deposition of an ARC/ashing layer (e.g., AlN layer) and then into the intermediate transfer chamber 722. The second robot 730 moves the substrate from the intermediate transfer chamber 722 into the PVD chamber 732, 734, 736 or 738. A hardmask layer (e.g., hardmask layer 205 in FIG. 2) may be formed on the substrate in the PVD Chamber 732, 734, 736 or 738. The second robot 730 then picks up the substrate from the PVD Chamber 732 and transfers the substrate into the intermediate transfer and cooling chamber 724. The first robot 710 moves the substrate to the load lock 706B, so that the subsequent lithographic operations can be performed on the substrate external to the cluster tool 700.

In some process flows, it may be desirable for the hardmask layer containing substrate to be further processed in the cluster tool 700, or more typically be processed in a separate cluster tool that is configured similarly to the cluster tool shown in FIG. 7. In either case, the patterned resist containing substrate is placed in the load lock chamber 706A. The first robot 710 then loads the substrate into the thermal processing chamber 716. The substrate is exposed to a degassing process in the thermal processing chamber 716. The first robot 710 then picks the substrate from the thermal processing chamber 712 and transfers the substrate through the transfer chamber 724 to the second robot 730 and into the etch chamber 714 for etching of the hardmask and thereafter ashing of the substrate. The cluster tool 700 may move the substrate from the etch chamber 714 to the thermal treatment chamber 716 for a subsequent degassing. This process may repeat itself until a complete pattern is formed in the hardmask layer and the substrate is placed in the etch chamber 714 for etching the underlying layer.

FIGS. 4A to 4M show a sequence of fabrication of a plurality of intersecting trenches 492, 492 into the optically tunable "memorization" or hardmask layer 423. In FIG. 4A an underlying substrate, into which a pattern will ultimately be etched from the pattern of the hardmask, is shown. In FIG. 4b, an antireflective coating (ARC), such as a TiN ARC layer 422, is formed over the substrate 421. Thereafter, an optically tunable hardmask layer 423 as described herein is deposited over the arc layer 422 as seen in FIG. 4C. A resist layer 424, having optical properties matched to the hardmask layer, is formed thereon such as by spin coating to yield the structure as shown in FIG. 4D.

Referring now to FIG. 4E, after exposing the resist layer in a pattern through a mask (not shown) in a lithography

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process, the resist layer **424** is developed, leaving behind resist layer **424** with features **430** therein, separated by resist having a width or "critical dimension" between features of **432**. FIG. 4F shows the top view of the resist covered substrate **421** having the features **430** developed therein. Thereafter, the substrate **421** is exposed to an reactive ion etch environment, wherein an etch gas selective to etching the underlying hardmask material is introduced and energized in a plasma, and the substrate or substrate holder is biased to directionally etch openings **440** at least partially inwardly of the surface of the hardmask layer **423** copying the pattern developed in the resist. As shown in FIG. 4G, the resulting structure has openings **440** separated by a wall of the hardmask layer **423** maintaining the critical dimension **432** formed in the resist layer **424** during exposure and developing thereof. Thereafter, the remaining portion of the resist overlaying the patterned hardmask layer **403** is stripped in an O₂ or H₂/O₂ remote plasma, to ash (strip) the resist without significantly or meaningfully affecting the optical properties of the underlying hardmask layer **403**, resulting in the profile shown in FIG. 4H.

Referring now to FIG. 4I, the partially etched hardmask layer has a second photo-resist layer **441** formed thereon by spin coating or the like, wherein the second resist also is optically matched to the optically tuned hardmask layer **403**. Thereafter, as shown in FIG. 4K, trench features are exposed and developed in the resist layer **441**, wherein the trench features **460** extend orthogonally to the trench features **430** in the first resist layer **441**. In FIGS. 4J to 4L, the substrate is rotated 90 degrees compared to the previous Figures. Thereafter, the underlying hardmask layer **423** is etched through the resist layer **441** features to provide the trenches **480** as shown in FIGS. 4L and 4M.

By using an optically tunable hardmask, multiple near nanopatterns, in the FIG. 4 trenches, may be formed in consecutive patterning steps on the same hardmask layer, by only stripping off of the used resist, cleaning the surface, and applying a new resist, thus significantly reducing the time, cost, and complexity of multi-patterning of a hardmask.

Although descriptions in this disclosure relate to the methods for matching the refractive index (n) and extinction coefficient (K) of a hardmask to a photo-resist, the same means for matching these properties can be applied to other material layers as well. For instance, depositing a matching material can be considered for use as ARC films, Passivation films or buffer films. PVD films are differentiated for their high purity, high density and capability for low temperature deposition. Use of this invention could change the patterning materials and film stacks currently in use. Thus, the methods for forming a hardmask may be applied to a variety of layers and materials undergoing a lithographic operation.

While the foregoing is directed to embodiments of the present invention, other and further embodiments of the invention may be devised without departing from the basic scope thereof, and the scope thereof is determined by the claims that follow:

What is claimed is:

1. A method for forming a hardmask on a film stack, comprising:
sputtering a material comprising silicon from a target disposed in a chamber onto a surface of a substrate; and delivering a flow of a process gas while sputtering the material from the target, wherein the process gas comprises oxygen and nitrogen, and
wherein a ratio of oxygen to nitrogen in the process gas is adjusted so that optical properties of the sputtered material has substantially similar values as the optical

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properties of a photo-resist layer, which is to be disposed on a surface of the sputtered material, at an intended lithography exposure wavelength.

2. The method of claim 1, wherein the optical properties include a refractive index and an extinction coefficient.

3. The method of claim 2, wherein the exposure wavelength is 193 nm.

4. The method of claim 1, wherein the process gas further comprises carbon.

5. The method of claim 2, wherein the optical properties of the sputtered material are substantially different than the optical properties of the photoresist at a pattern alignment wavelength.

6. The method of claim 1, wherein the sputtered material comprises silicon, nitrogen, oxygen and hydrogen, wherein a concentration of hydrogen at the surface of the sputtered material is less than an average hydrogen concentration through a thickness of the sputtered material, or a concentration of nitrogen at the surface of the sputtered material is less than an average nitrogen concentration through the thickness of the sputtered material.

7. The method of claim 1, further comprising:
biasing an electrode coupled to a portion of the chamber to increase the hydrophobicity of the surface of the substrate.

8. The method of claim 1, further comprising depositing the photo-resist layer directly on the surface of the sputtered material.

9. The method of claim 8, wherein the optical property includes both a refractive index and an extinction coefficient, and the deposited photo-resist layer has both the refractive index between 1.5 and 1.8 and the extinction coefficient is between 0.00 and 0.12 at a wavelength of 193 nm.

10. The method of claim 1, further comprising:
depositing the photo-resist layer directly on the surface of the sputtered material, wherein the photo-resist layer comprises a first photo-resist layer;
patterning the sputtered material using the first photo-resist layer;
depositing a second photo-resist layer directly on a surface of the patterned sputtered material, wherein an optical property of the second photo-resist layer has a substantially equivalent value as the optical property of the patterned sputtered material at the second photo-resist layer's lithography exposure wavelength; and
patterning the sputtered material using the second photo-resist layer.

11. The method of claim 1, further comprising:
depositing the photo-resist layer directly on the surface of the sputtered material;
performing a lithography operation on the photo-resist layer to form a pattern in the photo-resist layer;
etching the formed pattern in the sputtered material; and
removing the photo-resist layer, wherein removing the photo-resist layer comprises ashing the photo-resist layer using a remote O₂ plasma or a remote H₂/N₂ plasma.

12. A method for forming a hardmask on a film stack, comprising:

sputtering a material comprising silicon from a target disposed in a chamber onto a surface of a substrate; and delivering a flow of a process gas while sputtering the material from the target, wherein the process gas comprises oxygen and nitrogen, and
wherein a ratio of oxygen to nitrogen in the process gas is adjusted so that optical properties of the sputtered material has substantially similar values as the optical

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properties of a photo-resist layer, which is to be disposed on a surface of the sputtered material, at an intended lithography exposure wavelength, wherein the process gas comprises a gas that is selected from a group consisting of argon, helium, neon, krypton, 5 xenon, nitrogen, forming gas, ammonia, oxygen, hydrogen, water, and a gas that comprises carbon.

13. The method of claim **12**, wherein the process gas comprises methane (CH_4), carbon monoxide (CO), or carbon dioxide (CO_2).

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